

PubMed Nucleotide Protein Genome Structure PMC Taxonomy OMIM Books

Search for

Limits Preview/Index History Clipboard Details

About Entrez

Show:

Text Version

1: Biochim Biophys Acta 1998 Aug 14;1373 (1):119-30

Entrez PubMed

Overview
Help | FAQ
Tutorial
New/Noteworthy
E-Utilities

PubMed
Services
Journals
Database
MeSH Database
Single Citation
Matcher
Batch Citation
Matcher
Clinical Queries
LinkOut
Cubby

Related
Resources
Order
Documents
NLM Gateway
TOXNET
Consumer Health
Clinical Alerts
ClinicalTrials.gov
PubMed Central

Privacy Policy

Cholesterol-dependent generation of a unique amyloid beta-protein from apically missorted amyloid precursor protein in MDCK cells.

Mizuno T, Haass C, Michikawa M, Yanagisawa K.

Department of Dementia Research, National Institute for Longevity Sciences, Gengo 36-3, Morioka, Obu 474, Japan.

To investigate the implications of altered sorting of the beta-amyloid precursor protein (betaAPP) in the abnormal generation of amyloid beta-protein (Abeta), we characterized Abeta secreted from Madin-Darby canine kidney (MDCK) cells which had been stably transfected with a cDNA encoding the human beta-amyloid precursor protein (betaAPP695) with a 42 amino acid residue truncation at the carboxyl terminus (DeltaC). In DeltaC MDCK cells, the intracellular sorting of betaAPP is substantially altered to the apical surface. We detected an accumulation of a unique Abeta species in the apical compartment of DeltaC MDCK cell cultures. This unique Abeta was immunoprecipitated with 4G8 (a monoclonal antibody specific for Abeta17-24) and detected as a smear on Western blots, but was not immunoprecipitated with BAN50 (a monoclonal antibody raised against Abeta1-16). Interestingly, however, this Abeta species was readily immunoprecipitated with BAN50 upon treatment with

formic acid. Furthermore, incubation of the DeltaC MDCK cells with compactin, an inhibitor of de novo cholesterol synthesis, or with filipin, a cholesterol-binding drug, resulted in marked changes in the characteristics of this Abeta species as follows: first, the Abeta was not observed as a smear on Western blots and second, the Abeta was immunoprecipitated with BAN50. The present results strongly suggest that an Abeta with unique molecular characteristics is generated from the missorted betaAPP in vivo in a cholesterol-dependent manner.

PMID: 9733943 [PubMed - indexed for MEDLINE]

Show:

[Write to the Help Desk](#)

[NCBI](#) | [NLM](#) | [NIH](#)

[Department of Health & Human Services](#)

[Freedom of Information Act](#) | [Disclaimer](#)

May 2 2003 16:34:23



National
Library
of Medicine 

PubMed Nucleotide Protein Genome Structure PMC Taxonomy OMIM [Book](#)
Search for

Limits [Preview/Index](#) [History](#) [Clipboard](#) [Details](#)

[About Entrez](#)

Show:

[Text Version](#)

1: Biochem Biophys Res Commun 1996 Mar [Related Articles](#)
27;220(3):710-8

[Links](#)

ELSEVIER SCIENCE
FULL-TEXT ARTICLE

Nuclear and cytoplasmic localization of the beta-amyloid peptide (1-43) in transfected 293 cells.

Johnstone EM, Babbey LE, Stephenson D, Paul DC, Santerre RF, Clemens JA, Williams DC, Little SP.

Central Nervous System/GI/GU/Molecular Biology
Research Division, Lilly Research Laboratories, Eli Lilly
and Company, Indianapolis, Indiana 46285, USA.

Cultures of transformed human embryonic kidney 293 cells were transiently transfected with minigene constructs coding for the Abeta peptide (1-43). The Abeta minigene used in this study consisted of exons 16 and 17 of the amyloid precursor protein gene, including the 6000+ bp intronic region. Two of the constructs used in this study, human amyloid precursor protein (APP) promoter-driven Abeta minigene and BK virus enhancer/adenovirus major late promoter-driven Abeta minigene, did not contain a signal peptide sequence, whereas the third, human APP promoter-signal peptide Abeta minigene did not contain the human APP signal sequence. The resulting Abeta products were detected by immune precipitation, using 10D5 antibody and Western blot analysis, using R1280 antisera, as SDS stable oligomers in cell lysates of cells containing all three constructs or in culture media when produced by the signal peptide construct. Evaluation of the cells by immunocytochemistry using conventional and

Related
Resources
Order
Documents
NLM Gateway
TOXNET
Consumer Health
Clinical Alerts
[ClinicalTrials.gov](#)
PubMed Central

[Privacy Policy](#)

transmission electron microscopy indicated that the cells transfected with constructs without the signal peptide accumulated immunoreactive Abeta primarily in the nucleus.

PMID: 8607830 [PubMed - indexed for MEDLINE]

Display Abstract Show: 20 Sort Send to Text

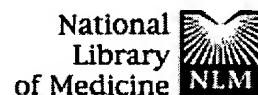
[Write to the Help Desk](#)

[NCBI](#) | [NLM](#) | [NIH](#)

[Department of Health & Human Services](#)

[Freedom of Information Act](#) | [Disclaimer](#)

May 2 2003 16:34:21



PubMed Nucleotide Protein Genome Structure PMC Taxonomy OMIM Books

Search for

Limits Preview/Index History Clipboard Details

About Entrez

[Text Version](#)

[Entrez PubMed](#)

[Overview](#)
[Help | FAQ](#)
[Tutorial](#)
[New/Noteworthy](#)
[E-Utilities](#)

[PubMed](#)
[Services](#)
[Journals](#)
[Database](#)
[MeSH Database](#)
[Single Citation](#)
[Matcher](#)
[Batch Citation](#)
[Matcher](#)
[Clinical Queries](#)
[LinkOut](#)
[Cubby](#)

[Related Resources](#)
[Order](#)
[Documents](#)
[NLM Gateway](#)
[TOXNET](#)
[Consumer Health](#)
[Clinical Alerts](#)
[ClinicalTrials.gov](#)
[PubMed Central](#)

[Privacy Policy](#)

Show: Sort Text

1: Biochemistry 1998 Oct 20;37
 (42):14958-65

[Related Articles](#)

[Links](#)



Processing of the Alzheimer's disease amyloid precursor protein in *Pichia pastoris*: immunodetection of alpha-, beta-, and gamma-secretase products.

Le Brocque D, Henry A, Cappai R, Li QX, Tanner JE, Galatis D, Gray C, Holmes S, Underwood JR, Beyreuther K, Masters CL, Evin G.

Department of Pathology, The University of Melbourne, Parkville, Victoria, Australia.

betaA4 (Abeta) amyloid peptide, a major component of Alzheimer's disease (AD) plaques, is a proteolytic product of the amyloid precursor protein (APP). Endoproteases, termed beta- and gamma-secretase, release respectively the N- and C-termini of the peptide. APP default secretion involves cleavage within the betaA4 domain by alpha-secretase. To study the conservation of APP processing in lower eukaryotes, the yeast *Pichia pastoris* was transfected with human APP695 cDNA. In addition to the full-length integral transmembrane protein found in the cell lysate, soluble/secreted APP (sAPP) was detected in the culture medium. Most sAPP comprised the N-terminal moiety of betaA4 and corresponds to sAPPalpha, the product of alpha-secretase. The culture medium also contained minor secreted forms detected by a monoclonal antibody specific for sAPPbeta (the ectodomain released by beta-secretase

cleavage). Analysis of the cell lysates with specific antibodies also detected membrane-associated C-terminal fragments corresponding to the products of alpha and beta cleavages. Moreover, immunoprecipitation of the culture medium with three antibodies directed at distinct epitopes of the betaA4 domain yielded a 4 kDa product with the same electrophoretic mobility as betaA4 synthetic peptide. These results suggest that the alpha-, beta-, and gamma-secretase cleavages are conserved in yeast and that *P. pastoris* may offer an alternative to mammalian cells to identify the proteases involved in the generation of AD betaA4 amyloid.

PMID: 9778373 [PubMed - indexed for MEDLINE]

Show:

[Write to the Help Desk](#)

[NCBI](#) | [NLM](#) | [NIH](#)

[Department of Health & Human Services](#)

[Freedom of Information Act](#) | [Disclaimer](#)

May 2 2003 16:34:23



National
Library
of Medicine
NLM

PubMed Nucleotide Protein Genome Structure PMC Taxonomy OMIM

Search for

Limits Preview/Index History Clipboard Details

[About Entrez](#)

Show:

[Text Version](#)

1: Brain Res 1999 Mar 27;823(1-2):202-6 Related Articles, Links

[Entrez PubMed](#)

[Overview](#)
[Help | FAQ](#)
[Tutorial](#)
[New/Noteworthy](#)
[E-Utilities](#)

[PubMed](#)
[Services](#)
[Journals](#)
[Database](#)
[MeSH Database](#)
[Single Citation](#)
[Matcher](#)
[Batch Citation](#)
[Matcher](#)
[Clinical Queries](#)
[LinkOut](#)
[Cubby](#)

[Related Resources](#)
[Order](#)
[Documents](#)
[NLM Gateway](#)
[TOXNET](#)
[Consumer Health](#)
[Clinical Alerts](#)
[ClinicalTrials.gov](#)
[PubMed Central](#)

[Privacy Policy](#)

Immunohistochemical localization of amyloid beta-protein with amino-terminal aspartate in the cerebral cortex of patients with Alzheimer's disease.

Arai T, Akiyama H, Ikeda K, Kondo H, Mori H.

Department of Neuropathology, Tokyo Institute of Psychiatry, 2-1-8 Kamikitazawa, Setagaya-ku, Tokyo 156-8585, Japan. arai@prit.go.jp

We investigated immunohistochemically the localization of amyloid beta-protein (Abeta) with amino-terminal aspartate (N1[D]) in brains of patients with Alzheimer's disease, diffuse Lewy body disease and Down's syndrome. A monoclonal antibody, 4G8, which recognizes the middle portion of Abeta, was used as a reference antibody to label the total Abeta deposits. Double staining with anti-Abeta (N1[D]) and 4G8 revealed that Abeta deposits in the subiculum and the neocortical deep layers often lacked N1[D] immunoreactivity, indicating N-terminal truncation of Abeta in these deposits. Abeta deposits in the neocortical superficial layers and the presubiculum parvopyramidal layer always contained Abeta with N1[D]. Such regional as well as laminar differences in the distribution of Abeta beginning at N1[D] suggest that some local factors influence N-terminal processing of Abeta deposited in the brain. Copyright 1999 Elsevier Science B.V.

PMID: 10095028 [PubMed - indexed for MEDLINE]

Show:

[Write to the Help Desk](#)

[NCBI](#) | [NLM](#) | [NIH](#)

[Department of Health & Human Services](#)

[Freedom of Information Act](#) | [Disclaimer](#)

May 2 2003 16:34:23



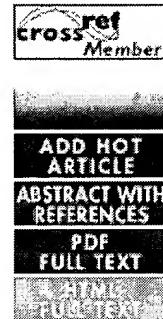
Online ISSN: 1098-1136 Print ISSN: 0894-1491

Glia

Volume 25, Issue 4, 1999. Pages: 324-331

Published Online: 8 Feb 1999

Copyright © 1999 Wiley-Liss, Inc.



Original Article

Occurrence of the diffuse amyloid β -protein (A β) deposits with numerous A β -containing glial cells in the cerebral cortex of patients with Alzheimer's disease

Haruhiko Akiyama ^{1*}, Hiroshi Mori ¹, Takaomi Saito ², Hiromi Kondo ¹, Kenji Ikeda ¹, Patrick L. McGeer ³

¹Tokyo Institute of Psychiatry, 2-1-8, Kamikitazawa, Setagaya-ku, Tokyo, Japan

²Laboratory for Proteolytic Neuroscience, RIKEN Brain Science Institute, Wako, Japan

³Kinsmen Laboratory of Neurological Research, University of British Columbia, Vancouver, British Columbia, Canada

email: Haruhiko Akiyama (akiyama@prit.go.jp)

* Correspondence to Haruhiko Akiyama, Tokyo Institute of Psychiatry, 2-1-8, Kamikitazawa, Setagaya-ku, Tokyo, 156-8585, Japan.

Funded by:

- Ministry of Education, Science and Culture of Japan; Grant Number: 08670743, 07264244
- the Naito Foundation

- Mitsui Foundation
- Jack Brown and Family Alzheimer's Disease Research Fund

Keywords

$\text{A}\beta$ 40; microglia; astrocytes; clearance; uptake; degradation

Abstract

Diffuse amyloid β -protein ($\text{A}\beta$) deposits with numerous glial cells containing C-terminal $\text{A}\beta$ fragments occur in the cerebral cortex of patients with Alzheimer's disease. By using a panel of antibodies specific for various epitopes in the $\text{A}\beta$ peptide, we have investigated the immunohistochemical nature of the diffuse $\text{A}\beta$ deposits. The extracellular material contains $\text{A}\beta$ with a C-terminus at residue valine⁴⁰ ($\text{A}\beta$ 40) as well as residues alanine⁴²/threonine⁴³ ($\text{A}\beta$ 42). The N-termini include aspartate¹, pyroglutamate³, and pyroglutamate¹¹, with pyroglutamate³ being dominant. Microglia and astrocytes in and around these deposits contain intensely staining granules. Most of these granules are negative for antibodies to the N-terminally located sequences of $\text{A}\beta$. These include 6E10 ($\text{A}\beta$ 1-17), 6F/3D ($\text{A}\beta$ 8-17), and the N-terminal antibodies specific to aspartate¹, pyroglutamate³, and pyroglutamate¹¹. The C-termini of intraglial $\text{A}\beta$ are comparable with those of the extracellular deposits. The microglia and astrocytes have quiescent morphology compared with those associated with senile plaques and other lesions such as ischemia. Complement activation in these deposits is not prominent and often below the sensitivity of immunohistochemical detection. Although factors which may cause this type of deposit remain unclear, lack of strong tissue responses suggests that these deposits are a very early stage of $\text{A}\beta$ deposition. They were found only inconsistently and were absent in a number of cases examined in this study. Further analysis of these deposits might provide important clues regarding the accumulation and clearance of $\text{A}\beta$ in Alzheimer's disease brain. GLIA 25:324-331, 1999. © 1999 Wiley-Liss, Inc.

Received: 13 April 1998; Accepted: 20 August 1998

References are available in the Enhanced Abstract

Additional Information

- Find other articles like this in Wiley InterScience.

- Find articles in Wiley InterScience written by any of the authors.
- Find other links for this article.

[\[Wiley InterScience Home Page\]](#) [\[Personal Home Page\]](#) [\[Journal Finder\]](#) [\[Book Finder\]](#)
[\[Search Wiley InterScience\]](#) [\[Reference Works\]](#) [\[Help\]](#) [\[Contact Us\]](#)[\[Logout\]](#)

Wiley InterScience is a member of **CrossRef**



Copyright © 1999-2003 by John Wiley & Sons, Inc. All rights reserved.